

Influenza-Associated Pediatric Deaths Case Report Form

FAX THIS FORM TO: COMMUNITY EPIDEMIOLOGY BRANCH AT (619) 515-6644

Form approved OMB No. 0920-0007

STATE USE O	STATE USE ONLY – DO NOT SEND INFORMATION IN THIS SECTION TO CDC					
Last Name:	First Name:		County:			
Address:	City:		State, Zip:			
Patient Demographics						
1. State: 2. Con	unty:	3. State ID:	4. CDC ID:			
5. Age: Days G. Date G	me of birth:/	/ 7.Sex: ☐ Male ☐ Female	[☐ Hispanic or Latino ☐ Not Hispanic or Latino ☐ Unknown		
9. Race: ☐ White ☐ Black ☐ Asian ☐ Native Hawaiian or Other Pacific Islander ☐ American Indian or Alaska Native ☐ Unknown						
Death Information						
10. Date of illness onset:// 11. Date of death:// to CDC? MM DD YYYY						
Test Type	Result			Specimen Collection Date		
☐ Commercial rapid diagnostic test	☐ Influenza A ☐ Influenza B ☐ Negative ☐ Influenza A/B (Not Distinguished)		//			
☐ Viral culture			/			
☐ Direct fluorescent antibody (DFA)	□ Influenza A □ Influenza B □ Negative □ Influenza A/B		//			
☐ Indirect fluorescent antibody (IFA)	□ Influenza A □ Influenza B □ Negative □ Influenza A/B		//			
☐ Enzyme immunoassay (EIA)	☐ Influenza A (Subtyping I☐ Influenza A (Unable To	Not Done) □ Influenza B Subtype) □ Influenza A (H1)	☐ Negative ☐ Influenza A (H3)	/		
□ RT-PCR	☐ Influenza A (Subtyping I☐ Influenza A (Unable To	Not Done) □ Influenza B Subtype) □ Influenza A (H1)	☐ Negative ☐ Influenza A (H3)	/		
☐ Immunohistochemistry (IHC)	☐ Influenza A	□ Influenza B	☐ Negative	//		



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Culture confirmation of INVASIVE bacterial pathogens				
14 a. Was a specimen collected for bacterial culture from a normally sterile site (e.g., blood, cerebrospinal fluid [CSF], tissue, or pleural fluid)?				
14 b. If yes, please indicate the site from which the specimen was obtained. □ Blood Date/_/_ □ Positive □ Negative □ Unknown □ Pleural fluid Date/_/_ □ Positive □ Negative □ Unknown □ CSF Date/_/_ □ Positive □ Negative □ Unknown □ Other Date//_ □ Positive □ Negative □ Unknown □ Unknown				
14 c. What was the result of the bacterial culture?		☐ Positive ☐ Negative ☐ Unknown		
14 d. If positive, please check the organism cultured.				
☐ Streptococcus pneumoniae	☐ Staphylococcus aureus, methicillin sensitive	□ Neisseria meningitidis (serogroup, if known):		
☐ Haemophilus influenzae type b	☐ Staphylococcus aureus, methicillin resistant (MRSA)	☐ Group A streptococcus		
☐ Haemophilus influenzae not-type b	☐ Staphylococcus aureus, sensitivity not done	☐ Other invasive bacteria:		
Culture confirmation of bacter	rial pathogens from NON-STERILE SIT	ΓES		
14 e. Were other respiratory specimens collected for bacterial culture (e.g., sputum, ET tube aspirate)? ☐ Yes ☐ No ☐ Unknown				
14 f. If yes, please indicate the site from which the specimen was obtained. □ Sputum □ Date _/_/ □ Positive □ Negative □ Unknown □ ET tube □ Date _/_/ □ Positive □ Negative □ Unknown □ Other □ Date _/_/ □ Positive □ Negative □ Unknown □ Unknown				
14 g. What was the result of the bacteri	al culture?	☐ Positive ☐ Negative ☐ Unknown		
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☐ Haemophilus influenzae type b	☐ Staphylococcus aureus, methicillin resistant (MRSA)	☐ Group A streptococcus		
☐ <i>Haemophilus influenzae</i> not-type b	☐ Staphylococcus aureus, sensitivity not done	☐ Other bacteria:		
Medical Care				
15. Did the patient receive medical care for this illness before admission to the hospital or death if outside the hospital? ☐ Yes* ☐ No ☐ Unknown				
16. If YES*, indicate level(s) of care received (check all that apply): □ Outpatient clinic □ ER □ Inpatient ward □ ICU				

☐ Yes

 \square No

 \square Unknown

17. Did the patient require mechanical ventilation?



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Clinical Diagnoses and Complications						
18 a. Did complications occur during the acute illness: ☐ Yes ☐ No ☐ Unknown						
18 b. If yes, check all complications that occurred during the acute illness:						
□ Pneumonia (Chest X-Ray confirmed) □ Acute Respiratory Disease Syndrome (ARDS) □ Croup □ Seizures						
□ Bronchiolitis □ Encephalopathy/encephalitis □ Reye syndrome □ Shock □ Sepsis						
□ Another viral co-infection: □ Other: □						
19 a. Did the child have any medical conditions that existed before the start of the acute illness: ☐ Yes ☐ No ☐ Unknown						
19 b. If yes, check all medical conditions that existed before the start of the acute illness:						
☐ Moderate to severe developmental delay ☐ Hemaglobinopathy (e.g. sickle cell disease) ☐ Asthma/ reactive airway disease						
☐ Diabetes mellitus ☐ History of febrile seizures ☐ Seizure disorder ☐ Cystic fibrosis						
□ Cardiac disease (specify) □ Skin or soft tissue infection						
☐ Chronic pulmonary disease (specify) ☐ Immunosuppressive condition (specify)						
☐ Metabolic disorder (specify) ☐ Neuromuscular disorder (including cerebral palsy) (specify)						
☐ Pregnant (specify gestational age) weeks ☐ Other (specify)						
Medication and Therapy History						
20 a. Was the patient receiving any of the following therapies in the 7 days prior to illness onset or after illness onset? (check all that apply) 20 b. Was the patient receiving any of the following therapies prior to illness onset? (check all that apply)						
☐ Aspirin or aspirincontaining products ☐ NSAID or NSAID or NSAID or notation products ☐ Antibiotic therapy or radiation specify therapy specify therapy specify therapy specify therapy or radiation specification therapy or radiation specification therapy specification therapy or radiation specification therapy specification t						
Influenza vaccine history						
21. Did the patient receive any influenza vaccine during the current season (before illness) ☐ Yes* ☐ No ☐ Unknown						
22. If YES* , please specify influenza vaccine received before illness onset: □ Trivalent inactivated influenza vaccine (TIV) [injected] □ Live-attenuated influenza vaccine (LAIV) [nasal spray] □ Unknown						
23. If YES*, how many doses did the patient receive and what was the timing of each dose? (Enter vaccination dates if available)						
□ 1 dose □ <14 days prior to illness onset ONLY □ ≥14 days prior to illness onset Date dose given:// MM DD YYYY						
24. Did the patient receive any influenza vaccine in previous seasons? ☐ Yes ☐ No ☐ Unknown						
Submitted By: Date: / / Phone No.: () MM DD YYYY E-mail Address:						

Public reporting burden of this collection of information is estimated to average 20 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC/ATSDR Reports Clearance Officer; 1600 Clifton Road NE, MS E-11, Atlanta, Georgia 30333; ATTN: PRA (0920-0007).